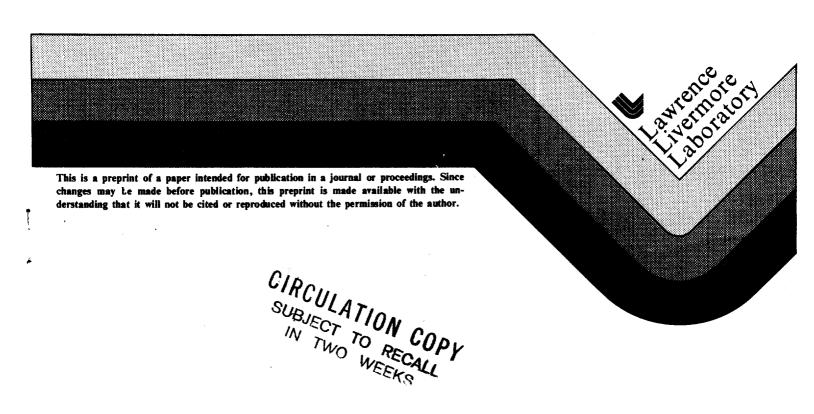
THE TANDEM MIRROR REACTOR AS A SYNTHETIC FUEL PRODUCER

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Introduction

We believe that fusion should and must play as strong a role in the production of fuels and chemical feedstocks as it is expected to play in the production of electricity. The role, in fact, may be even stronger since the production of fuel in the form of hydrogen, hydrocarbons, and their derivatives useful for transportation, industrial processes, or for residential and commercial use or the production of chemicals based on hydrogen or hydrocarbons is three times as high in the United States as is the use of energy for the production of electricity.

We further believe that fuel production is not inimical to electricity production from fusion but is complementary to it and strengthens the base of the entire fusion program.

We report on a scoping design of a fusion reactor based on tandem mirror physics coupled to thermochemical processes for the production of hydrogen.

Flow of Energy in the U.S.

The reason for coupling synfuels to fusion, in terms of a U.S. energy problem, is that the principal shortfall in the future United States energy flow will continue to be in liquid and gaseous fuels as it currently is and not in electrical power. In the time period for fusion's emergence, it is likely that the demand for electrical energy will be outweighed by the demand for energy in the form of portable fuels by a ratio of three to one. Incorporating synthetic fuel production in the fusion program both broadens the fusion program's base and is responsive to a national need.

Figure 1 illustrates the U.S. energy flow picture for 1979. The future overall demand would be expected to increase by perhaps 5% per year on the average if the past is any guide.

*Work performed under the auspices of the U.S. Department of Energy by the Lawrence Livermore National Laboratory under contract number W-7405-ENG-48.

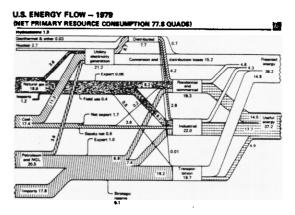


Figure 1

The division of energy flow should not be expected to change materially.

Even though our fusion reactor may be 20 or 30 years away from commercial fruition, now is an appropriate time to begin to plan and study the fusion/synfuel tie. An interesting way to begin is with the tandem mirror reactor (TMR) which, as we shall see, has some distinct advantages over other energy producers for this application.

The Technical Choice of the TMR

The fusion reactor as an energy source was critical to our study, and we investigated how well fusion might produce synthetic fuels, as well as what influence the fuel production would have on basic reactor design.

It can be stated that the main influence synfuel production had on reactor design had to do with blanket modules, those units surrounding the plasma that convert the neutrons' kinetic energy to thermal energy and in the case of the D-T cycle also produce the tritium part of the fueling needs by neutron-lithium reactions. Other engineering elements of the reactor remain substantially the same. The physics may be slightly easier due to the basic size of a fusion reactor for synfuels compared to one for electrical production. The synfuel plant is larger by perhaps a factor of two and thus the

reactor Q can become higher.

TMR Design Parameters - Thermochemical Hydrogen vs. Electrical Production

Table 1 provides a comparison between a conceptual reactor design for fuel production and one for electrical production.

TABLE 1

Example Design Parameters for the Tandem Mirror Reactor For Synfuel Production or Electrical Production

		<u>Synfuel</u>	Elec.
Pusion Power	(10f _£)	5680	3500
Thermal Power	()ef _{th})	4945	3360
Pirst Wall Loading	() () () () () () () () () () () () () (2.0	2.6
HCMM Power Delivered	(900)	330	260
Nautral Beam Power Delivered			
Central Cell	(997)	0.0	0
Berrier Cell	(997)	85	58
Plug	(180)	8.5	48
Central Cell Length	(m)	260	125
Central Cell First Wall Redius	(m)	1.4	1.7
Global Reactor Q-Value		13.3	9.6
Prest Ac	(16fe)	0 .	1000
3c	(T)	~2	~2
a _p	(T)	~12	~12

Assembly, Disassembly and Accessibility

A primary technical reason for choosing the tandem mirror was its highly favorable reactor configuration. The configuration not only allows the design of relatively simple blanket modules that are the principal source of the process heat but puts them together into a workable package.

The sequence of figures that follow, Figs. 2A, B, C, D, E, F, G; illustrates how the blanket modules, tailored for synfuel, thermochemical cycle use may be assembled into a highly manageable, serviceable, accessible reactor for producing energy.

The Operating Principle of the Cauldron Blanket Module

A cross-sectional view of the cauldron module is illustrated in Figure 3. Notice the module's resemblance to a pool boiler. It is, however, substantially more complicated than a pool boiler due to geometric effects, due to the exponential energy generation in the fluid contained within the module and, last but not least, due to MHD effects on the convective mixing of the two liquid metals, lithium and

sodium, we have chosen to use in the pool. The two liquids in this cauldron module act as both the neutron moderator and heat transfer fluid, absorbing energy in direct proportion to the energy input and transferring it by latent heat of vaporization of the sodium to a heat exchanger in the dome of the vessel. The

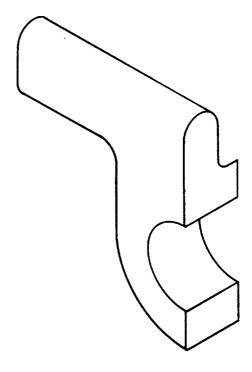


Figure 2A A single cauldron blanket module

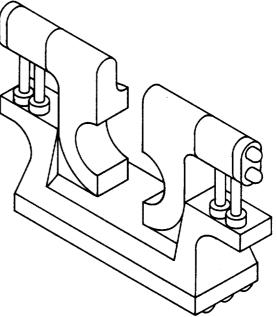
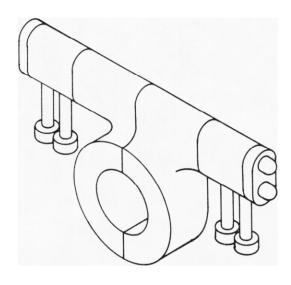


Figure 2B A retracted pair of cauldron modules or an assembly



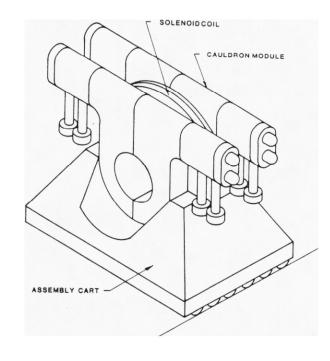


Figure 2C Two modules ready for assembly with a central cell solenoid

Figure 2D A unit cell or its assembly cart

CROSS SECTION THRU 2 UNIT CELLS

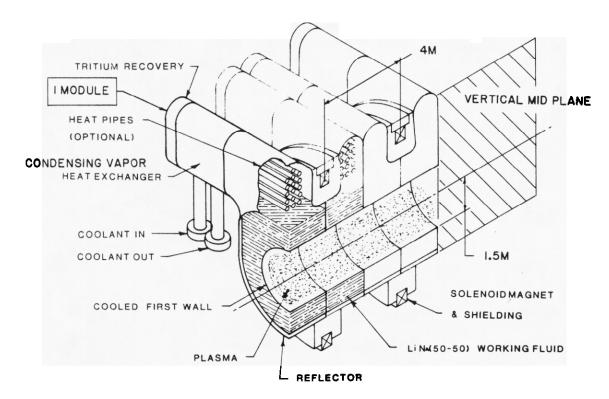


Figure 2E

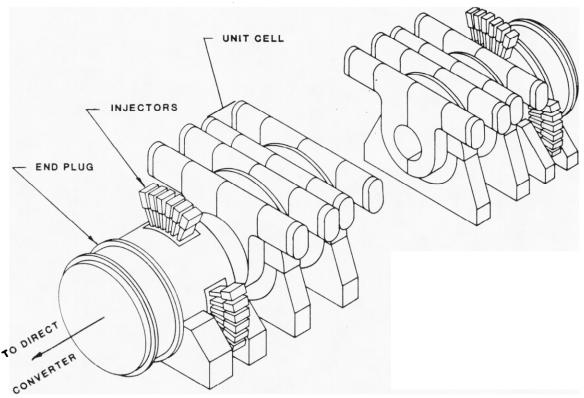


Figure 2F Unit cells installed in reactor

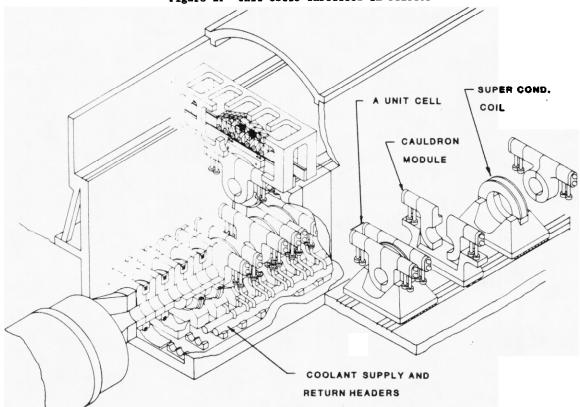


Figure 2G A part of the main reactor showing states of assembly or disassembly

NAVAPOR

NAVAPOR

NAVAPOR

PLASMA

PLASMA

PLASMA

PLASMA

DEPLIS ATM

TODA

TODA

REFLECTOR

SOLENOID COIL

ENERGY
DEPLIN

BLANKET

100

DEPLIN

BLANKET

COND. VAPOR HEAT EXCHANGER

Figure 3

lithium performs the function of tritium breeding. In the dome the condensing vapor heat exchanger (CVHX) transfers the thermal energy out of the module to various chemical processors located some distance from the reactor.

The sodium preferentially vaporizes, leaving behind the lithium in the liquid state to do its neutron moderating, tritium-producing function. The sodium vapor, traveling at vapor velocities roughly 8-10 m/s at 1200 K, condenses on the heat exchanger tubes in the dome, yields energy and returns as liquid droplets to the pool, thus completing the cycle. We have selected the LiNa mixture for our studies but also considered potassium. The two fluids are miscible and tritium production with a 50-50 mix is greater than 1. The neutronics of the potassium is a little less favorable although the Lik can run cooler for the same vapor velocity. LiRb or LiCs are other mixes or compounds that may be of interest.

The Tandem Mirror Reactor and Its Energy Sources

The tandem (TMR) is a steady state, driven, fusion device. It operates on the deuterium-tritium cycle. Energy from the reactor is produced in two primary forms as evidenced by equation 1.

Deuterium + Tritium Energetic Neutron + charged particle

The first energy form, the kinetic energy of the neutron, is captured in the moderating blanket surrounding the reacting plasma and thermal energy is produced. The cylindrical plasma is physically located in the TMR's central cell in a zone that is about 200 meters long for the synfuel production application. The plasma is contained within the central cell by the retarding action of the mirror end cells whose electrostatic potential causes the deuterium and tritium ions to be reflected back and forth sufficiently long so that some of them react with one another and fuse. Figure 4 illustrates this potential and the magnetic fields associated with the tandem.

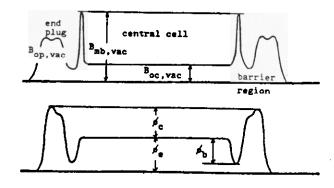


Figure 4

The second energy form is that contained in the charged alpha particle. On forming, the alpha begins to lose some of its 3.5 Mev energy by heating the plasma through a series of collisions with electrons and ions and finally. at some degraded energy, leaves the system (the central cell) through the ends as do the deuterium and tritium ions that did not react. All the alpha particle energy may now be accounted for as the sum of its residual energy plus the enhanced energy of the exiting ions. This energy is recovered in a direct convertor located beyond the end cells. The direct convertor produces two forms of energy, an electrical dc component which we use in our point design to drive the reactor, and a thermal component which, for our thermochemical cycle, furnishes energy for process chemistry. The neutrons, as they are moderated in the blanket, produce some additional energy by exothermic

TABLE 2

Partitioning the Energy for Process Chemistry

At the blanket region it is convenient for structural and thermodynamic reasons to divide all of the blanket energy into two sources of supply usable for the process chemistry. The first supply source is the cooled first wall zone, characterized by modest (700 K) energy levels since the first wall serves as the structural container for the fluid in the module. The second zone is where high temperature, high-quality heat (1200 K) is produced for the main part of the thermochemical processes.

The Thermochemical Cycles

The TMR is used as an energy source to produce fuel (H₂) via three candidate thermochemical cycles.

°	Sulfur-Indine Cycle: (General Atomic)
	2 H ₂ 0 + SO ₂ + I ₂ AQUEOUS H ₂ SO ₄ + 2 HI _x
	2 HI _x -573K x I ₂ + H ₂
	H ₂ SO ₄ 11.44K H ₂ O + SO ₂ + 1/2 O ₂
•	Sulfur Cycle (part electrochemical): (Westinghouse)
	2 H ₂ 0 + SO ₂ AQUEOUS H ₂ + H ₂ SO ₄
	H_2SO ₄ HIGH T > H_20 + SO ₂ + 1/2 O ₂
•	Sulfur-Browine Cycle (part electrochemical): (ISPRA)
	2 H ₂ 0 + SO ₂ + Br ₂ AQUEOUS 320-370K → H ₂ SO ₄ + 2 HBr
	2 HB _r AQUECUS Rr ₂ + H ₂
	$H_2SO_4 \xrightarrow{1000-1100K} H_2O + SO_2 1/2 O_2$

These three cycles have been demonstrated on a laboratory scale.

The requirements for energy, thermal and electrical, for these three cycles are illustrated in table 2.

The Reactor Energy Balance, Thermal and Electric

For the thermochemical cycles, as Table 2 suggests, we are interested in limited electrical power production but primarily in the production of process heat and in the use of the available thermal energy out of the fusion reactor. Two regions of the tandem mirror fusion reactor provide this process heat: (1) the blanket and (2) the direct convertor. We will also use any surplus electrical energy out of the direct convertor over and above that

Electrical and Thermal Requirements for Various
Cycles Based on HTGR Heat Sources

Thermo-	Thermal	71	700000	Thermal energy us	ed to generate
chemical	met.	linet		electricity or shaft work	
Cycle					
			,	ļ	
		High	Interned.	Electrolytic	Process
		7000	Temp	Demand	Shaft Work
General Atomic	474	244	518	0	259
Sulfur Indine		1250 E	843 K		
	 				
- Westinghouse	47%	234	204	57%	0
Sulfur cycle		1280 K	1106 E		
					
LEPNA	46%	27%	52%	218	٥
Hack-13		1063 E	pelox		
			773 K		

which will be needed to satisfy circulating (driving) power requirements for the reactor itself.

The reactor energy balance, as it relates to the thermochemical cycles, is illustrated in Figure 5. We have established the following quantities which may be used with this energy balance.

AM =	The blanket thermal energy.	(1)
$B(1-\eta_{dc}) =$	The direct convertor thermal energy.	(2)
Bη _{dc} =	The direct convertor electrical energy.	(3)
c =	Thermal energy contribution from injected neutrals.	(4)
where, for th	ne D-T cycle:	
where, for th	0.8 P _{inj} $\eta_{A} \widehat{Q}$ (The neutron energy fraction)	(5)
В =	(The neutron energy fraction) $0.2 P_{inj} \eta_A Q + P_{inj} \eta_A$ (The alpha or charged particle	(6)
c -	fraction) P _{inj} (1 - η _A) (The fraction of injected neutrals that don't get ionize	(7) d)

The definitions of terms needed for Figure 5 are:

$\eta_{ extsf{A}}$	ionization efficiency.
M	blanket energy multiplication (from exothermic reactions).
η_{i}	injector efficiency.
$\eta_{ ext{inj}} \ \eta_{ ext{dc}}$	direct convertor efficiency (con- verting charged particle kinetic
	energy to electrical).
$\eta_{ m TH1}$	thermal convertor efficiency (con- verting thermal energy to electrical.
$\eta_{_{TH2}}$	 thermal convertor efficiency.
$\eta_{ ext{TH2}}$	power required for pumps, refrigera- tion, and other fusion plant needs
ô	= fusion power produced/power injected

POWER BALANCE MIRROR REACTOR
WITH DIRECT CONVERSION PROCESS
HEAT PRODUCTION FOR SYNFUELS

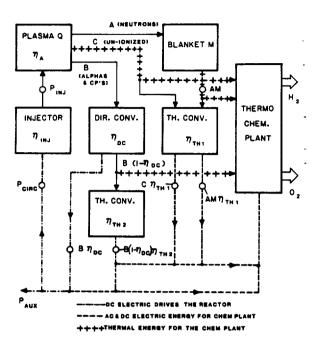


Figure 5

The value of $P_{\mbox{\scriptsize aux}}$ is taken as a fraction, f, of the thermal and dc energy before conversion

$$P_{aux} = f (B + AM + C)$$
 (8)

By definition

$$P_{circ} = P_{inj}/\eta_{inj}$$
 (9)

The Uniqueness of the TMR

There is a uniqueness associated with the tandem mirror reactor open-ended physics geometry that increases the usefulness of this energy source over other contemporary energy producers such as solar energy concentrators, tokamaks, and high temperature gas cooled fission reactors (HTGR) for the production of synthetic fuels. This uniqueness may be noted in the energy balance figure and has to do with the fact that in the TMR open-ended mirror geometry there is a direct convertor which produces dc power that may be used in thermochemical cycles that have an electrical demand and/or an electrochemical step. The result of using the direct convertor is an improvement in overall efficiency as schematically illustrated in Figs. 6A and 6B.

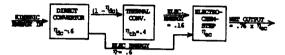
The Influence of Q on the TMR's Uniqueness

To fully recognize the TMR as a unique energy source for synfuel (H₂) production, it will be stipulated that the dc electrical energy



The STGR, the tokensk, solar energy conventrators <u>must convert</u> thermal energy to electrical energy at an efficiency $\eta_{\rm th}$ to drive the electrolysis step or meet other electrical demands of the thermochemical cycle.

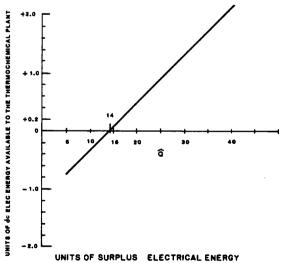
Figure 6A HTGR, Tokamak or solar energy concentrator energy sources



The tandom mirror reactor can drive the electrolysis step <u>directly</u> or provide other electrical demands <u>directly</u> using the reactor's do electrical output.

Figure 6B The tandem mirror reactor energy source

component Bndc of the direct convertor will be used, in the first place, to satisfy the circulating power and the auxiliary needs of the reactor itself. This means simply that the dc component of the direct convertor drives the reactor and the ancillary equipment. No other energy input is required. There will be a specific value of Q at which this demand is exactly satisfied. For higher Q values, there will then be a surplus of dc electrical energy that will then be used by the thermochemical plant. No electrical energy is sold. For lesser values of Q, some thermal energy from the reactor blanket or from the thermal part of the direct convertor would have to be converted to electrical energy to help drive the reactor. Figure 7 is a plot of how this electrical surplus



UNITS OF SURPLUS ELECTRICAL ENERGY VS Q. TANDEM MIRROR REACTOR

P_{SURPLUS}* [P_{RC}-(P_{AUX}+P_{CIRC})] P_{INJ}* 1.0

Figure 7

or deficiency, Pdc-(Paux + Pcirc), varies as a function of Q. The representative set of data for this plot are given in Table 3:

TABLE 3

Representative Quantities for the Energy Balance

Pini	injection power	= 1.0 Unit
$\eta_{\mathbf{a}}^{\mathtt{P_{inj}}}$	ionization efficiency	= 0.95
M	blanket energy multiplication	1.20
η_{m}	thermal efficiency	= 0.40
$\eta_{ ext{TH}} \ \eta_{ ext{inj}}$	injector efficiency	= 0.60
f	fraction of thermal and dc	
	energy before conversion	= 0.02
$\eta_{ exttt{dc}}$	direct convertor efficiency	= 0.55

It may be seen from Figure 7 that when the Q value is about 14 or higher, there begins to be a dc electrical component left over from the reactor that can be used to drive such things as electrolysis cells, pumps, motors, etc., in the synfuel plant.

Relative to the total energy that the reactor has created for synfuel purposes, the fraction that is dc tends to a limit R, where R is defined as surplus dc expressed in units of equivalent thermal energy divided by the total energy available to the process chemistry in thermal units. Expressing surplus dc in thermal units is somewhat of an artifice but is useful in illustrating limits. This limit is indicated in Figure 8 where we see that as Q gets larger and larger, the dc percentage of the reactor's output available for process chemistry tends to a limit of about 13%.

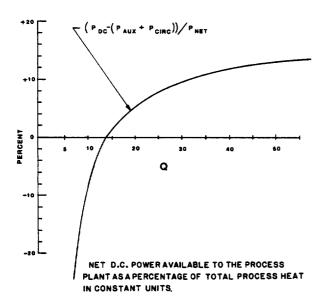


Figure 8

Comments on the Tandem as an Energy Source

From Figures 7 and 8 some conclusions can be reached.

- At a Q value ~14 the TMR dc electrical output is just large enough to feedback and drive the reactor, leaving the thermal fraction of the direct convertor and the blanket thermal energy available for process chemistry.
- As Q values exceed 14 there is some surplus dc electrical power available for process chemistry.
- As Q increases the direct current electrical energy that is available for process chemistry tends to a limit of about 13% of the plant useful output.
- 4) The availability of this direct current electrical energy from the TMR is an asset that other energy producing machines do not have. The HTGR, the tokamak, the LWR, the FBR -- all of these machines must go through the thermal conversion step to produce this electricity at a pensity that is directly proportional to the thermal efficiency. The tandem begins to avoids the thermal step when values of Q exceed 14.

Looking to the Future - The TMR and the DD or D-He3 Cycle

When we consider only the D-T fuel cycle, there is a limit for the charged particle energy out of the TMR that cannot exceed 20% of the total energy output i.e., 3.5 Mev alphas (14.1 Mev neutrons + 3.5 Mev alphas).

If a D-D fuel cycle were to be considered the picture changes significantly. With the D-D cycle it is possible to have approximately 50% of the raw energy output of the TMR in charged particle form and convertible to electricity directly. It is interesting to compare the two cycles (the DT and the DD) for their overall plant efficiency potential and also to compare the TMR with the tokamak, the HTGR or a solar concentrator under these circumstances. This comparison is shown below in Fig. 9.

Planning Ahead

We have included comments on the tandem D-D fuel cycle as a closure to this paper. We are of the opinion that the D-D cycle, although difficult from a physics standpoint, may be significantly superior to D-T from an engineering/technology viewpoint. The economics may be in question because of poorer reaction cross-section. However, the environmental/political influences and pressures that will

o TMRDBH D-T
$$\eta_{\rm mat}$$
 = .8 $(\eta_{\rm th})$ + .2 $\left[(\eta_{\rm th})$ + $(1-\eta_{\rm dr})\right]\eta_{\rm th}$ $\eta_{\rm mat}$ = .32 + 15 = 476

o THEOREM D-D
$$\eta_{\text{met}}$$
 = .5 (η_{ch}) + .5 $\left[(\eta_{\text{de}}) + (1 - \eta_{\text{de}})\right] \eta_{\text{th}}$
 $\eta_{\text{met}} \simeq .20 + .38 = 584$

o TOWNER D-T or D-D,
the STGR, or a
SOLAR CONCENTRATOR.

Fig. 9 Net Efficiency potential of different energy sources.

inevitably be brought to bear on fusion's acceptability to the community cannot be ignored. The inexhaustible energy advantages of deuterium fuel over tritium fuel are also important. We consider the D-D cycle or other advanced cycles as a necessary backup for the D-T cycle. We recommend that the fusion community begin to seriously assess these cycles using today's physic's base and avoiding the pitfall that our perceptions ten years ago indicated that D-T was difficult enough and we didn't need another order of magnitude problem. Since then physics and engineering progress has moved us several orders of magnitude closer to success. In the meantime political and social pressures against "things nuclear" have increased an order of magnitude. We must always view fusion from all areas: technical, political, social, institutional, environmental so that as we grow larger in national importance we retain our advocates.

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